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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/808,184	03/24/2004	Raghavan Rajagopalan	1486.1:H US (073979.68)	4580
27805 THOMPSON H	7590 10/25/201 IINE L.L.P.	EXAMINER		
Intellectual Prop P.O. BOX 8801	perty Group	PACKARD, BENJAMIN J		
DAYTON, OH		ART UNIT	PAPER NUMBER	
			1612	
			MAIL DATE	DELIVERY MODE
			10/25/2010	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)		
10/808,184	RAJAGOPALAN ET	AL.	
Examiner	Art Unit		
Benjamin Packard	1612		

	Denjamin Fackard	1012	
The MAILING DATE of this communication appe	ars on the cover sheet with the o	correspondence add	ress
THE REPLY FILED <u>18 October 2010</u> FAILS TO PLACE THIS A	PPLICATION IN CONDITION FO	R ALLOWANCE.	
1. The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following rapplication in condition for allowance; (2) a Notice of Appe for Continued Examination (RCE) in compliance with 37 C periods:	replies: (1) an amendment, affidavi al (with appeal fee) in compliance	t, or other evidence, wwith 37 CFR 41.31; or	which places the r (3) a Request
a) The period for reply expiresmonths from the mailing	date of the final rejection.		
b) The period for reply expires on: (1) the mailing date of this Adno event, however, will the statutory period for reply expire la Examiner Note: If box 1 is checked, check either box (a) or (I MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f	ter than SIX MONTHS from the mailing b). ONLY CHECK BOX (b) WHEN THE).	g date of the final rejection FIRST REPLY WAS FI	on. LED WITHIN TWO
Extensions of time may be obtained under 37 CFR 1.136(a). The date of have been filed is the date for purposes of determining the period of extremely an extra transfer of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	ension and the corresponding amount hortened statutory period for reply origi	of the fee. The appropria nally set in the final Offic	ate extension fee be action; or (2) as
2. ☐ The Notice of Appeal was filed on . A brief in compl	iance with 37 CFR 41.37 must be	filed within two month	s of the date of
filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed wi AMENDMENTS	sion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	
3. The proposed amendment(s) filed after a final rejection, b			cause
(a) They raise new issues that would require further cor		ΓE below);	
(b) They raise the issue of new matter (see NOTE below	**	d	ha iaawaa fan
(c) ☐ They are not deemed to place the application in bett appeal; and/or	er form for appeal by materially red	ducing or simplifying ti	ne issues for
(d) ☐ They present additional claims without canceling a c	orresponding number of finally reje	ected claims.	
NOTE: (See 37 CFR 1.116 and 41.33(a)).			
4. The amendments are not in compliance with 37 CFR 1.12	1. See attached Notice of Non-Co	mpliant Amendment (PTOL-324).
5. Applicant's reply has overcome the following rejection(s):			
 Newly proposed or amended claim(s) would be allenon-allowable claim(s). 	owable if submitted in a separate,	timely filed amendmer	nt canceling the
7. For purposes of appeal, the proposed amendment(s): a) [how the new or amended claims would be rejected is prov The status of the claim(s) is (or will be) as follows:		l be entered and an e	xplanation of
Claim(s) allowed:			
Claim(s) objected to: Claim(s) rejected: <u>11,12 and 21-27</u> .			
Claim(s) withdrawn from consideration: <u>13-20</u> .			
AFFIDAVIT OR OTHER EVIDENCE			
 The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e). 			
9. The affidavit or other evidence filed after the date of filing a entered because the affidavit or other evidence failed to or showing a good and sufficient reasons why it is necessary	vercome <u>all</u> rejections under appea	al and/or appellant fail	s to provide a
10. The affidavit or other evidence is entered. An explanation	n of the status of the claims after e	ntry is below or attach	ed.
REQUEST FOR RECONSIDERATION/OTHER 11. The request for reconsideration has been considered but	does NOT place the application in	condition for allowan	ce because:
See Continuation Sheet. 12 Note the attached Information Disclosure Statement(s) (DTO/SR/08) Donor No/o)		
12. ☐ Note the attached Information <i>Disclosure Statement</i>(s). (13. ☐ Other:	r I O/SD/U0/ Maper INO(S).		
/Frederick Krass/	/Benjamin Packard/		
Supervisory Patent Examiner, Art Unit 1612	Examiner, Art Unit 1612		

Continuation of 11. does NOT place the application in condition for allowance because: Claim Rejections - 35 USC § 103 Claims 11-12 and 21-27 stand rejected under 35 U.S.C. 103(a) as being unpatentable over Cuttitta et al (US 5,460,801) in view of Pandurangi et al (J. Org. Chem. 1998, 63, 9019-9030).

Applicants assert the instantly claimed method is directed to and recites "a phototherapeutic procedure" which results in photoexcitation of the Ar and N3 components, which phototreat the target tissue. In contrast, Applicants note Cuttitta's method binds to bombesin receptors but does not provide motivation or suggestion for a phototherapeutic procedure. Applicants assert Pandurangi doesn't cure this where it teaches perfluoroaryl azies "capable of complexing transitional metal" and producing photolibale chelating agents in vitro which are used "for the development of highly efficient CH insertion molecular probes". Such use is different than Applicants type 1 phototherapy where the aryl nitrene portion of the compound actually treats the target tissues.

Examiner disagrees. First, Cuttitta was not cited for the phototherapy component, but for the teaching that receptor specific administration were known in the art, specifically compounds used to bond to bombesin receptors.

Second, the step of the photoactivated aryl nitrene group to cause "cell death", i.e. type 1 phototherapy, is not in the instant claims. Although the claims are interpreted in light of the specification, limitations from the specification are not read into the claims. See In re Van Geuns, 988 F.2d 1181, 26 USPQ2d 1057 (Fed. Cir. 1993). Here, the claims are directed instead to simply "treat the target tissue". Where "treatment" is not defined in the instant specification, it is broadly interpreted as including any phase of the treatment, from diagnostics to killing of the cells. Here, treatment is interpreted to include a compound made obvious which has a nitrene for photoactivation in order to provide CH insertion into the adjacent molecules, thereby acting as a photolabeling agent. Note, the aryl nitrenes of Pandurangi et al were tested in vitro for the ability to photolabel human serum albumin, which provides a reasonable expectation that it will work similarly with other biological components.

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